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Jc713 U.S. PTO



09/22/99

Case No. 5050/582

PATENT APPLICATION TRANSMITTAL LETTER

To the Assistant Commissioner for Patents:

Transmitted herewith for filing is the patent application of: Randall L. Schlesinger and Mathew Rahimi for : MEDICAL DIAGNOSTIC
ULTRASOUND CATHETER WITH DIELECTRIC ISOLATION. Enclosed are:

- ☒ 2 sheet(s) of drawings, 14 pages of application (including title page), and the following Appendices : _____.
- ☒ Declaration.
- ☒ Power of Attorney.
- ☐ Verified statement to establish small entity status under 37 CFR §§ 1.9 and 1.27.
- ☒ Assignment transmittal letter and Assignment of the invention to : Acuson Corporation.
- ☒ Check for \$40.00 Recordal of Assignment Fee.

Jc678 U.S. PTO
09/401632
09/22/99

Claims as Filed	Col. 1	Col. 2
For	No. Filed	No. Extra
Basic Fee		
Total Claims	37-20	17
Indep. Claims	5-3	2
Multiple Dependent Claims Present		

If the difference in col. 1 is less than zero,
enter "0" in col. 2.

Small Entity	
Rate	Fee
	\$ 380
x\$9=	\$
x\$39=	\$
0+\$130=	\$0
Total	\$

Other Than Small Entity	
Rate	Fee
	\$ 760
17x\$18=	\$306.00
2x\$78=	\$156.00
0+\$260=	\$0
Total	\$1222.00

Please charge my Deposit Account No. 23-1925 in the amount of \$: _____. A duplicate copy of this sheet is enclosed.

A check in the amount of \$: 1,222.00 to cover the filing fee is enclosed.

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Date

9/24/99

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"Express Mail" Mailing Label No.

September 22, 1999

Date of Deposit

Our Case No. 5050/582

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE
APPLICATION FOR UNITED STATES LETTERS PATENT

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MATHEW RAHIMI

TITLE:

MEDICAL DIAGNOSTIC
ULTRASOUND CATHETER WITH
DIELECTRIC ISOLATION

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MEDICAL DIAGNOSTIC ULTRASOUND CATHETER WITH DIELECTRIC ISOLATION

BACKGROUND

5 This invention relates to medical diagnostic ultrasound catheters. In particular, different materials for constructing medical diagnostic ultrasound catheters provide improvements in dielectric isolation.

10 Catheters for use with ultrasound systems include transducers for converting electrical energy to acoustic energy and vice versa. Images of a region of a patient adjacent to the transducer are generated. In order to provide accurate imaging, a window or lens is integrated with the catheter adjacent to the transducer. The remainder of the exterior of the catheter generally comprises polyurethane or other biologically acceptable material. In order to provide torque transmission and flexibility, an internal metal braid is provided.

15 For safe inter-cardiac use of a catheter, the catheter must dielectrically withstand 3,000 volts for 60 seconds between the external surface of the catheter and any internal conductors or electrically active components when internal voltages are greater than 50 volts. Additionally, any leakage current must be less than 50 microamps when 264 volts are applied to the internal conductors. Alternating current capacitive coupling causes a voltage to be applied to the metal braid when voltage is applied to the internal conductors. The capacitive coupling decreases the dielectric withstand strength of the catheter and requires a thicker coating material, such as the polyurethane. However, smaller catheters are generally desired.

20 Components are often thermally fused with the shaft of the catheter. Since
25 the fusing is thermal, portions of the metal braid, such as exposed end wires, may migrate towards the outside surface of the catheter. A fused catheter more likely will not pass the safety requirements due to migration of the metal braid, resulting in wasted production. The metal braid may migrate to the surface, allowing potential corrosion.

These safety requirements apply to the entire catheter, including the tip area with the transducer. The adjacent lens or window material must satisfy the safety requirements. In order to provide the proper dielectric withstand, the thickness of the lens or window material is increased. However, the required thickness may be considerable. The greater the thickness, the larger the catheter or the less room for transducer components, such as desired additional transducer elements and associated conductors.

To avoid increasing the thickness of the shaft or lens of the catheter, each conductor may be isolated with a transformer. Such isolation requires a different dielectric withstand standard for safety, such as a 300 volt standard. For ultrasound imaging with high resolution, multiple conductors are used, such as 64 to 128 conductors. Isolating each conductor with a transformer is cost prohibitive.

BRIEF SUMMARY

The present invention is defined by the following claims, and nothing in this section should be taken as a limitation on those claims. By way of introduction, the preferred embodiment described below includes a medical diagnostic ultrasound catheter with materials for improved dielectric withstand strength. In one aspect, the catheter includes a braid of non-conductive material. The non-conductive braid reduces the capacitive coupling effects and allows smaller catheters or a greater number of conductors. The non-conductive braid provides both compressive and tensile strength to transmit the torque applied to the catheter. The non-conductive braid also allows fusing of components while decreasing the risk of defective manufacture.

In another aspect, a dielectric film, such as a polyester film, is positioned between the transducer and any lens or window. The dielectric film allows thinner lenses to be used, allowing smaller catheters or larger transducers. The dielectric film may also increase the sensitivity of the transducer to acoustic energy. The dielectric film prevents the lens material from filling kerfs in the transducer, which may eliminate the need for filling the kerfs of the transducer.

In one embodiment, a medical diagnostic ultrasound catheter for imaging from within a body is provided. The catheter includes a conductor and a shaft surrounding the conductor. A non-conductive braid or insert is connected with the shaft. An ultrasound transducer is connected with the shaft and electrically connects with the conductor.

In other embodiments, a medical diagnostic ultrasound catheter for imaging from within a body includes a dielectric film adjacent to an ultrasound transducer. The ultrasound transducer is connected with a shaft. A lens is adjacent to the ultrasound transducer. In other embodiments, the lens is optional and the dielectric film is between a portion of the shaft and the ultrasound transducer.

Further aspects and advantages of the preferred embodiments are described below in conjunction with the detailed description of the preferred embodiments.

BRIEF DESCRIPTION OF SEVERAL VIEWS OF THE DRAWINGS

FIG. 1 is a perspective view, in partial cutaway, of one preferred embodiment of a portion of a medical diagnostic ultrasound catheter.

FIG. 2 is a perspective view of one preferred embodiment of a transducer and associated conductor for use in the catheter of FIG. 1.

FIG. 3 is a cross-sectional view of one preferred embodiment of a tip portion of the catheter of FIG. 1.

FIG. 4 is a cross-sectional view of one preferred embodiment of a body portion of the catheter of FIG. 1.

FIG. 5 is a perspective view of a preferred embodiment of a medical catheter.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The preferred embodiments of the medical diagnostic ultrasound catheter include one or both of (1) a non-conductive insert for providing torque transmission and (2) a dielectric film extending around at least a portion of a transducer. The non-conductive insert described below prevents or reduces capacitive coupling, allowing for thinner catheter walls. The dielectric film

described below allows for less thick lenses adjacent to the transducer while still providing sufficient dielectric withstand for the 3,000 volts safety requirement.

Referring to FIGS. 1 and 5, a medical diagnostic ultrasound catheter is generally shown at 10. The catheter 10 includes a body portion 12 and a tip portion 14. A handle and control portion 15 (FIG. 5) connects with the body portion 12. The tip portion 14 includes a transducer 16 and associated conductors 18. The conductors also pass through the body portion 12. The catheter 10, including the body portion 12 and tip portion 14, comprises a shaft 20. While the shaft 20 shown is generally linear, shafts with angles, interruptions and other configurations may be used. In alternative embodiments, the transducer 16 is positioned in other locations different than the tip portion 14 of the catheter 10.

Referring to FIG. 2, the transducer 16 and associated conductors 18 are shown separately from the catheter. The transducer 16 comprises a piezoelectric material. The piezoelectric material is diced into a plurality of transducer elements 22 separated by kerfs 24. In one preferred embodiment, 64 transducer elements 22 are provided. In alternative embodiments more or fewer transducer elements may be used. As shown in FIG. 2, the transducer elements 22 are aligned linearly along a flat surface, but in alternative embodiments one and a half or two dimensional arrays of the elements 22 may be used. In yet other alternative embodiments, some or all of the transducer elements 22 are curved and/or not flat.

The upper surface 26 of the transducer 16 comprises an emitting face. A backing block 28 minimizes the acoustical energy transmitted from the transducer 16 in a direction other than through the emitting face 26.

Each transducer element 22 or a group of transducer elements 22 is electrically connected with a respective conductor 18. In one preferred embodiment, the transducer elements 22 are connected to the conductors 18 through a folded flexible circuit 30. The conductors 18 preferably comprise metalized copper conductors on a polyimide substrate (flex circuit). Each circuit has 18 conductors, including ground conductors, where four circuits are stacked to form a bundle.

FIG. 3, shows a cross-section of the tip portion 14 of the catheter 10. The tip portion includes the transducer 16, the backing block 28 and the folded flexible circuit 30. These components are within the shaft 20. The emitting face 26 of the transducer 16 is positioned adjacent a lens section 32 of the shaft 20. The lens section 32 preferably comprises Pebax®, such as Pebax® 3533 manufactured by Elf Autochem, but other polymers, silicones or materials may be used. The lens 32 is shaped to provide no focus or some focus, such as a physical elevation focus for the transducer 16 and is of a material that does not interfere with the transmission of acoustic energy. As used herein, lens includes a window or other structure on the shaft 20 for transmitting acoustic energy between the patient and the transducer 16, including windows that do or do not focus the acoustic waves.

In one preferred embodiment, a dielectric film 34 extends around the circumference and ends of the transducer 16 and associated components, including the backing block 28 and flexible circuit 30. The dielectric film 34 comprises any film providing dielectric insulation. Preferably, a polyester film is used, such as a Mylar® Type C polyester film manufacturing by DuPont. In alternative embodiments, the dielectric film 34 extends only between the emitting face 26 and the shaft 20. In yet other alternative embodiments, the dielectric film is positioned around only part of the transducer 16, backing block 28, flexible circuit 30, and/or conductors 18. The film may be a sheet or a tube.

The dielectric film 34 insulates the outer surface of the catheter 10 from the conductive components associated with the transducer 16. Preferably, the dielectric film 34 is thin enough to not degrade the acoustic signal but thick enough to provide high dielectric withstand strength. For example, the dielectric film 34 may have a thickness of about six microns. A high dielectric withstand strength may allow the thickness of the lens material 34 to be reduced. At six microns, the Mylar® Type C manufactured by DuPont provides a dielectric strength of 2,000 volts. Other thicknesses can be used, including greater or lesser thicknesses.

In one preferred embodiment, the dielectric film 34 comprises a tape. For example, a backing is peeled off of the dielectric film 34 to expose an adhesive.

The adhesive is then used to adhere to the transducer 16 and associated components.

The dielectric film 34 provides dielectric insulation, allowing for a larger transducer 16 or a smaller catheter 10. The dielectric film 34 also prevents the lens or shaft material from entering the kerfs 24 of the transducer 16 during manufacture, avoiding a kerf-filling step.

FIG. 4, shows a cross-sectional view of the body portion 12 of the catheter 10. The body portion 12 includes the shaft 20, lumens 36 for control lines to guide the catheter 10, and a lumen 38 for housing the conductors 18. More or fewer lumens 36 for control lines may be provided. In alternative embodiments, more than one lumen 38 for housing the conductors 18 is provided.

The shaft 20 comprises a polyurethane or other material. Preferably, the shaft 20 comprises Pebax[®], such as Pebax[®] 7233 manufactured by Elf Autochem. The lumens 36 and 38 are bonded or fused at each end of the catheter 10 to the shaft 20 or other components of the catheter 10. In this embodiment, the shaft 20 is hollow. In alternative embodiments, the shaft 20 is not hollow, or the control and/or conductor lumens 36 and/or 38 are fused at other locations within the catheter 10.

The shaft 20 also includes a non-conductive insert 40. The non-conductive insert preferably is configured as a braid embedded within the shaft 20 during manufacture of the shaft 20. The non-conductive insert 40 comprises any non-conductive material, such as plastic, glass, nylon or combinations thereof. In one preferred embodiment, a liquid crystal polymer, Vectron[®] manufactured by Hoechst-Celanese Acetate, is used. Preferably, the material used comprises a monofilament material. In alternative embodiments, multiple fibers may be used. For example, the non-conductive insert 40 comprises a linear strip, a tube, a spiral winding, a diamond weave or other braid structures using two or more filaments or groups of filaments. Single-stranded or multi-stranded braids may be used. Preferably, a 32 or 64 psi (crossing per inch) diamond weave braid of 2 or 3 mil monofilament is used. The non-conductive insert 40 provides flexibility while transmitting the torque appropriate for use of the catheter 10. For example, the

non-conductive insert 40 has a modular of elasticity greater than 4Mpsi, such as 10Mpsi and transmits torque along substantially the entire length of the shaft 20.

In one preferred embodiment, the tip portion 14 is thermally fused to the body portion 12 during manufacture. More flexibility may be provided in the catheter 10 if the non-conductive insert 40 does not extend throughout the length of the tip portion 14. The desired rigidity and torque control is provided by embedding the non-conductive insert 40 within the body portion 12.

By using the non-conductive insert 40, the shaft 20 is maintained free of electrically conducting materials. This may reduce the capacitive coupling effect caused by metallic braids. During the thermal fusing operation, portions of the non-conductive insert 40 may migrate closer to an exposed surface of the catheter 10. Such migration does not result in failure of the catheter 10 due to safety requirements. Where the non-conductive insert 40 migrates to the surface, corrosion and harm to the patient is prevented.

While the invention has been described above by reference to various embodiments, it will be understood that many changes and modifications can be made without departing from the scope of the invention. For example, a non-conductive insert may be used without the dielectric film. Likewise the dielectric film may be used without the non-conductive insert. The dielectric film may be used to wrap other electrically conductive parts, such as the conductors.

It is therefore intended that the foregoing detailed description be understood as an illustration of the presently preferred embodiments of the invention, and not as a definition of the invention. It is only the following claims, including all equivalents, that are intended to define this scope of the invention.

CLAIMS

What is claimed is:

1. A medical diagnostic ultrasound catheter for imaging from within a body, the catheter comprising:
5 a conductor;
 a shaft surrounding at least a portion of the conductor;
 a non-conductive braid connected with the shaft; and
 an ultrasound transducer connected with the shaft and electrically connected with the conductor.
- 10 2. The catheter of Claim 1 wherein the non-conductive braid comprises a braid of mono-filament material.
3. The catheter of Claim 1 wherein the non-conductive braid comprises mono-filament nylon material.
- 15 4. The catheter of Claim 1 wherein the non-conductive braid comprises a material selected from the group consisting of: glass, plastic, nylon and combinations thereof.
5. The catheter of Claim 1 wherein the non-conductive braid is embedded within the shaft.
- 20 6. The catheter of Claim 1 wherein the shaft is free of electrically conductive material.
7. The catheter of Claim 1 wherein the shaft comprises a tip portion fused to a body portion, the non-conductive braid connected with the body portion.
- 25 8. The catheter of Claim 1 wherein the non-conductive braid comprises a configuration selected from the group consisting of: spiral, diamond weave and combinations thereof.

9. The catheter of Claim 1 further comprising a dielectric film adjacent an emitting surface of the ultrasound transducer.

10. A medical diagnostic catheter for use within a body, the catheter comprising:

5 an electrical conductor;
a shaft surrounding at least a portion of the electrical conductor; and
a non-conductive insert for providing torque transmission connected with the shaft.

11. The catheter of Claim 10 wherein the non-conductive insert comprises a braid of mono-filament material.

12. The catheter of Claim 10 wherein the non-conductive insert is embedded within the shaft.

13. The catheter of Claim 10 wherein the shaft comprises a tip portion fused to a body portion, the non-conductive insert connected with the body portion.

14. The catheter of Claim 10 further comprising a dielectric film adjacent an emitting surface of a ultrasound transducer, the ultrasound transducer connected with the shaft and electrically connected with the conductor.

15. A medical diagnostic ultrasound catheter for imaging from within a body, the catheter comprising:

a shaft;
an ultrasound transducer connected with the shaft;
a lens adjacent the ultrasound transducer; and
a dielectric film adjacent the ultrasound transducer.

16. The catheter of Claim 15 wherein the dielectric film is positioned between the lens and the ultrasound transducer.

17. The catheter of Claim 15 wherein the dielectric film comprises a tape material.

18. The catheter of Claim 15 wherein the dielectric film comprises a polyester film.

5 19. The catheter of Claim 18 wherein the polyester film comprises Mylar.

20. The catheter of Claim 15 wherein the dielectric film comprises a film having a thickness less than 7 microns.

10 21. The catheter of Claim 15 wherein:
the shaft comprises a tip portion;
the ultrasound transducer connects with the tip portion; and
the dielectric film wraps around a circumference of the ultrasound transducer.

15 22. The catheter of Claim 15 wherein the dielectric film is positioned adjacent an emitting surface of the ultrasound transducer.

23. The catheter of Claim 15 further comprising a non-conductive braid connected with the shaft.

20 24. A medical diagnostic ultrasound catheter for imaging from within a body, the catheter comprising:

a shaft;
an ultrasound transducer connected within the shaft; and
a dielectric film positioned between a portion of the shaft and the ultrasound transducer.

25 25. The catheter of Claim 24 further comprising a lens adjacent the ultrasound transducer, wherein the dielectric film is positioned between the lens and the ultrasound transducer.

26. The catheter of Claim 24 wherein the dielectric film comprises a polyester film.

27. The catheter of Claim 24 wherein the dielectric film comprises a film having a thickness less than 7 microns.

5 28. The catheter of Claim 24 further comprising a non-conductive braid connected with the shaft.

29. A medical catheter for insertion into a body, the catheter comprising:

a shaft; and

10 a non-conductive braid connected with the shaft to transmit torque to the shaft.

30. The catheter of Claim 29 wherein the non-conductive braid comprises a braid of mono-filament material.

15 31. The catheter of Claim 29 wherein the non-conductive braid comprises mono-filament liquid crystal polymer material.

32. The catheter of Claim 29 wherein the non-conductive braid comprises a material selected from the group consisting of: glass, plastic, nylon and combinations thereof.

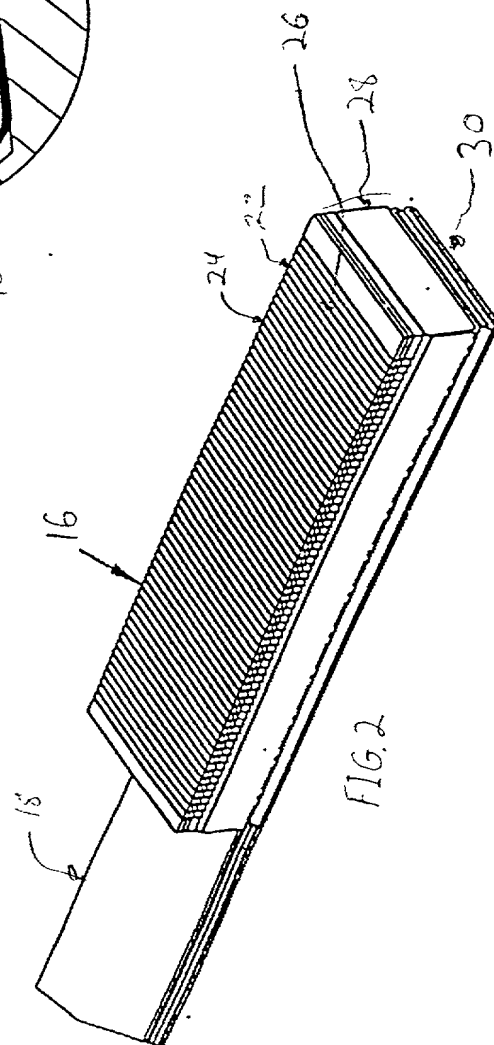
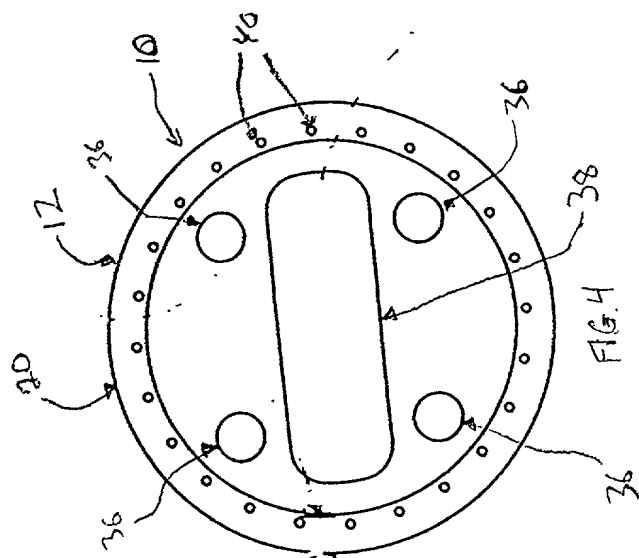
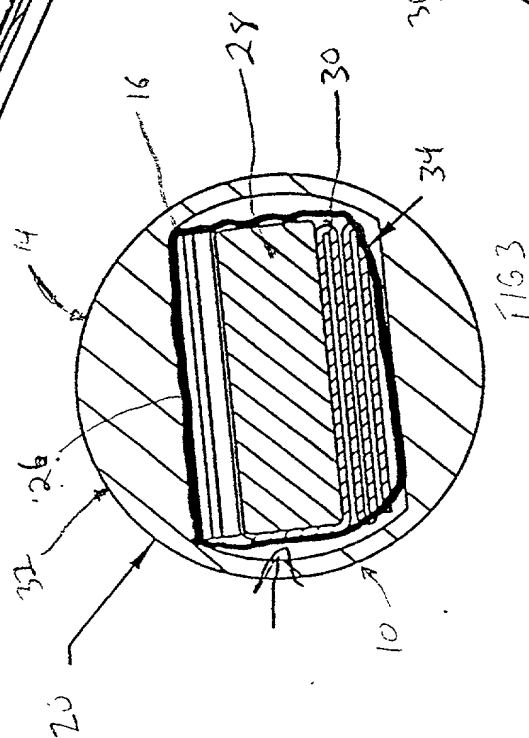
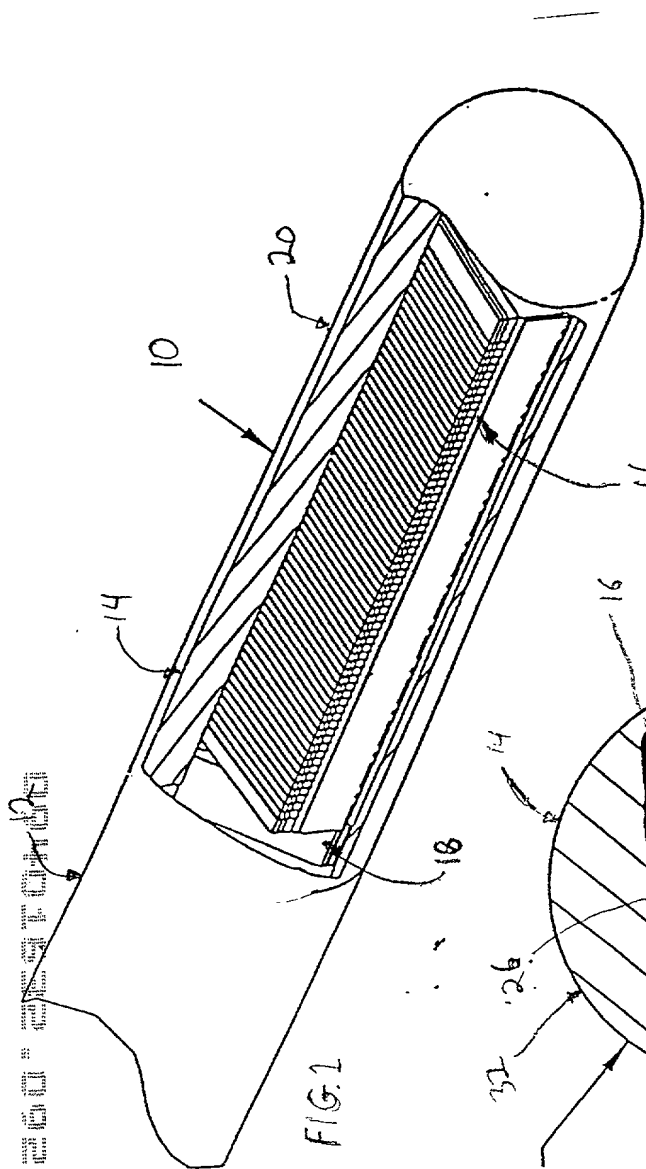
20 33. The catheter of Claim 29 wherein the non-conductive braid is embedded within the shaft.

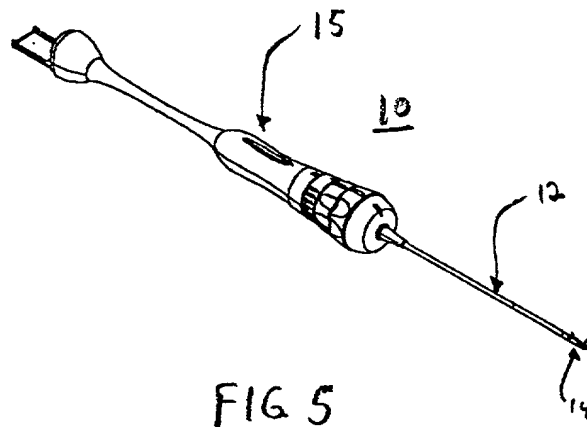
34. The catheter of Claim 29 further comprising:
a conductor surrounded by at least a portion of the shaft; and
an ultrasound transducer electrically connected to the conductor.

25 35. The catheter of Claim 29 further comprising a control within the shaft.

36. The catheter of Claim 1 wherein the non-conductive braid comprises liquid crystal polymer material.

37. The catheter of Claim 10 wherein the non-conductive insert comprises liquid crystal polymer material.





DECLARATION FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled MEDICAL DIAGNOSTIC ULTRASOUND CATHETER WITH DIELECTRIC ISOLATION, the specification of which:

- ☒ is attached hereto.
- ☐ was filed on _____ as Application Serial No. _____.
- ☐ and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the patentability as defined in Title 37, Code of Federal Regulations, § 1.56(a).

I hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s)Priority Claimed

_____	_____	_____	<input type="checkbox"/>	<input type="checkbox"/>
(Number)	(Country)	(Day/Month/Year Filed)	Yes	No

I hereby claim the benefit under 35 U.S.C. § 119(e) of any United States provisional application(s) listed below:

_____	_____
(Application Serial No.)	(Filing Date)

I hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application:

_____	_____	None
(Application Serial No.)	(Filing Date)	(Status-patented, pending, abandoned)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Inventor's Signature

Full name of sole or first inventor

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Date: 9/15/99

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Inventor(s): Randall L. Schlesinger and Mathew RahimiTitle: MEDICAL DIAGNOSTIC ULTRASOUND CATHETER WITH DIELECTRIC ISOLATION

**POWER OF ATTORNEY AND
37 C.F.R. §3.73(b) RIGHT OF ASSIGNEE TO PROSECUTE**

The specification of the above-identified patent application:



is attached hereto



was filed on _____, as application Serial No. _____

I hereby revoke all previously granted powers of attorney in the above-identified patent application and appoint the following attorneys to prosecute said patent application and to transact all business in the Patent and Trademark Office connected therewith:

William A. Webb (28,277)
Joseph F. Hetz (41,070)

Craig A. Summerfield (37,947)
Liza Toth (31,065)

Please address all correspondence and telephone calls to Craig A. Summerfield in care of:

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The undersigned hereby authorizes the U.S. attorneys named herein to accept and follow instructions from: Liza K. Toth, Esq., Registration No. 31,065 (415/694-5413) as to any action to be taken in the Patent and Trademark Office regarding this application without direct communication between the U.S. attorney and the undersigned. In the event of a change in the persons from whom instructions may be taken, the U.S. attorneys named herein will be so notified by the undersigned.

ACUSON CORPORATION, a Delaware Corporation, certifies that it is the assignee of the entire right, title and interest in the patent application identified above by virtue of either:

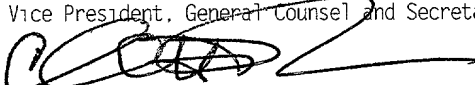
- A. ☒ An assignment from the inventor(s) of the patent application identified above, a copy of which is attached hereto.
- OR
- B. ☐ An assignment from the inventor(s) of the patent application identified above. The assignment was recorded in the Patent and Trademark Office at Reel _____, Frame _____.
- OR
- C. ☐ A chain of title from the inventor(s), of the patent application identified above, to the current assignee as shown below:
1. From: _____ To: _____
The document was recorded in the Patent and Trademark Office at Reel _____, Frame _____, or a copy thereof is attached.
 2. From: _____ To: _____
The document was recorded in the Patent and Trademark Office at Reel _____, Frame _____, or a copy thereof is attached.

☐ Additional documents in the chain of title are listed on a supplemental sheet.

The undersigned has reviewed the assignment or all the documents in the chain of title of the patent application identified above and, to the best of undersigned's knowledge and belief, title is in the assignee identified above.

The undersigned (whose title is supplied below) is empowered to act on behalf of the assignee.

I hereby declare that all statements made herein of my own knowledge are true, and that all statements made on information and belief are believed to be true; and further, that these statements are made with the knowledge that willful false statements, and the like so made, are punishable by fine or imprisonment, or both, under Section 1001, Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

Name: Charles H. DearbornTitle: Vice President, General Counsel and Secretary, ACUSON CORPORATIONSignature: Date: 9/17/99